

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 7:
A61N 1/365
A1
(11) International Publication Number: WO 00/57953
(43) International Publication Date: 5 October 2000 (05.10.00)

(21) International Application Number: PCT/SE00/00572

(22) International Filing Date: 23 March 2000 (23.03.00)

(30) Priority Data:
9901194-2
31 March 1999 (31.03.99)
SE

(71) Applicant (for all designated States except US): PACESETTER AB [SE/SE]; S-175 84 Järfälla (SE).

(72) Inventors; and

(75) Inventors/Applicants (for US only): MIN, Mart [EE/EE]; Söpruse pst. 188A-4, EE-Tallinn (EE). KINK, Andres [EE/EE]; Nabala tee 2A, Kiili, EE-75401 Harjumaa (EE). PARVE, Toomas [EE/EE]; Rännaku blvd. 3-7, EE-10917 Tallinn (EE).

(74) Common Representative: PACESETTER AB; Patent Department, Att: Sven Kalling, S-175 84 Järfälla (SE).

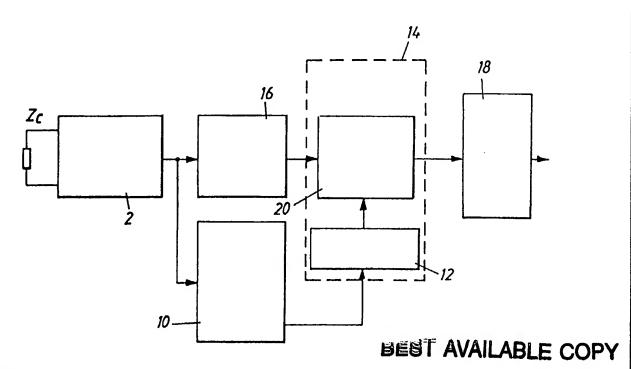
(81) Designated States: US, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).

Published

With international search report.

Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.

(54) Title: A RATE ADAPTIVE PACEMAKER



(57) Abstract

A rate adaptive pacemaker comprises a means (2) for determining the demand of the patient's organism, a pacing rate controlling means (16) for controlling the pacing rate in response to the patient's demand, and a pacing rate limiting means (20) for preventing the pacing rate from becoming too low. The pacing rate limiting means is adapted to limit the pacing rate downwards such that a first predetermined relation is satisfied between actual cardiac output (CO) and cardiac output (CO_{rest}) for the patient in rest conditions and a second predetermined relation is satisfied between actual stroke volume (SV) and rest stroke volume (SV_{rest}).

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia	
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia	
AT	Austria	FR	France	LÜ	Luxembourg	SN	Senegal	
ΑŪ	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland	
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad	
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo	
BB	Barbados	GH	Ghana	MG	Madagascar	ТJ	Tajikistan	
BE	Belgium	GN	Guinea	MK	The former Yugoslav	TM	Turkmenistan	
BF	Burkina Faso	GR	Greece		Republic of Macedonia	TR	Turkey	
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago	
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine	
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda	
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America	
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan	
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam	
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia	
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe	
CI	Côte d'Ivoire	KP	Democratic People's	NZ	New Zealand			
CM	Cameroon		Republic of Korea	PL	Poland			
CN	China	KR	Republic of Korea	PT	Portugal			
CU	Cuba	KZ	Kazakstan	RO	Romania			
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation			
DE	Germany	LI	Liechtenstein	SD	Sudan			
DK	Denmark	LK	Sri Lanka	SE	Sweden		•	
EE	Estonia	LR	Liberia	SG	Singapore			

WO 00/57953 PCT/SE00/00572

A RATE ADAPTIVE PACEMAKER Technical Field

The present invention relates to a rate adaptive pacemaker comprising a means for determining the demand of the patient's organism, a pacing rate controlling means for controlling the pacing rate in response to the patient's demand, and a pacing rate limiting means for preventing the pacing rate from becoming too low.

1

10 Background Art

5

15

20

The pacing rate of a rate adaptive pacemaker may become too low due to the physical demand of the patient's organism and heart. This may result in lack of oxygen supply to the myocardium. Under certain conditions the heart may not be able to fulfil the physiological needs of the patient's organism and heart if the pacing rate is not limited.

It is previously known to set a lower limit for the pacing rate. This limit value is normally determined from the patient's diagnosis and a constant or externally programmable limit can be set. Thus US-A-4,535,774 describes a stroke volume controlled pacemaker, in which the heart rate is permitted to range between prescribed minimum and maximum heart rate values. Further, in US-A-5,861,011 a pacemaker is disclosed having a system for determining the circadian rhythm by examining variations in the QT interval and adjusting the pacemaker night time setting of a lower rate limit to a lower value than the pacemaker daytime setting of the lower rate limit.

30

35

25

Thus, too low a pacing rate may cause too slow influx of blood enriched with oxygen. A prescribed suitable lower pacing rate limit avoids the slow influx of the fresh blood. At the same time this lower limit value should be low enough not to disturb a peaceful sleep. In that case the patient can feel more healthy in various everyday life conditions including peaceful sleeping.

The purpose of the present invention is to provide a rate adaptive pacemaker in which the pacing rate is prevented in a new way from becoming too low, such that the above discussed inconveniences for the patient are avoided.

Disclosure of the Invention

This purpose is obtained by a rate adaptive pacemaker according to claim 1.

10

15

5

Thus, by satisfying two predetermined relations the pacemaker according to the invention ensures a sufficient minimum energy supply to the patient's organism or body and at the same time the maximum value of the stroke volume is limited and these conditions are continuously automatically checked.

Preferred embodiments are set forth in the dependent claims.

According to an advantageous embodiment of the pacemaker according to the invention the first predetermined relation is

$$CO > CO_{rest}$$
 (1)

and said second predetermined relation is

$$(SV)/(SV_{rest}) < L$$
 (2)

where L denotes a predetermined constant > 1, preferably equal to a value between 1.2 and 1.5. In this way the actual cardiac output is ensured not to become lower than the rest state cardiac output CO_{rest} and the actual stroke volume is ensured to be less than a maximum allowed value equal to L × SV_{rest}, where L typically has a value between 1.2 and 1.5, depending on the health of the patient's myocardium. By satisfying both these conditions simultaneously a physiologically well founded heart work management at low work loads is ensured.

5

10

30

According to other advantageous embodiments of the pacemaker according to the invention the pacing rate limiting means includes a lower limit setting means for setting a lower limit value for the pacing rate, and a lower limit determining means for determining the relation between actual cardiac output (CO) and cardiac output (CO) and cardiac output (CO) and the relation between actual stroke volume (SV) and a rest stroke volume (SV) and calculating a lower pacing rate limit value from said relations for supply to said limit setting means, and said lower limit determining means includes a stroke volume measuring means for measuring actual stroke volume SV and comparison means for comparing measured actual stroke volume SV with stroke volume SV rest for the patient in rest conditions to ensure that the inequality

$$SV/SV_{rest} < L$$
 (3)

is satisfied and said lower limit determining means is adapted to calculate a lower pacing rate limit value from the equation

lower pacing rate limit =
$$HR_{rest} \cdot (SV_{rest}/SV)$$
 (4)

where HR_{rest} denotes the heart rate for the patient in rest conditions, provided that said inequality is satisfied. In this way the lower pacing rate limit is continuously automatically calculated and it may also happen that the lower pacing rate limit becomes lower than the typical heart rate HR_{rest} for rest conditions of the patient.

According to still another advantageous embodiment of the pacemaker according to the invention a bioimpedance measurement unit is provided to measure the cardiac bioimpedance as a function of time for determining therefrom actual cardiac output CO and actual stroke volume SV from the measured cardiac bioimpedance. In this way these parameters are obtained in an easy and reliable way from the time variation of the bioimpedance measured between a standard intracardiac

electrode and the housing of the pacemaker, when an excitation current proceeds from the electrode tip.

Brief Description of the Drawings

The invention will now be described more in detail with reference to the enclosed drawings on which figure 1 is a block diagram of an embodiment chosen as an example of the pacemaker according to the invention and figure 2 illustrates the principle of bioimpedance measurements between the tip of an intracardial electrode and the metal housing of the pacemaker.

Description of a Preferred Embodiment

To avoid that the current cardiac output CO

$$CO = SV \times HR \tag{5}$$

becomes lower than the rest state cardiac output CO_{rest} the pacing rate must be above a lower pacing rate limit given by

lower pacing rate limit =
$$(CO_{rest})/(SV)$$
 (6)

and since

$$20 CO_{rest} = HR_{rest} \times SV_{rest} (7)$$

lower pacing rate limit =
$$(HR_{rest}) \times (SV_{rest}/SV)$$
 (8)

In addition thereto the maximum value of the stroke volume must be limited, i.e.

$$SV < L \times SV_{rest}$$
 (9)

25

Thus, the following two conditions must be fulfilled simultaneously for insuring a physiologically well founded heart work management at low work loads.

Pacing rate limit >
$$(HR_{rest}) \times (SV_{rest}/SV)$$
 (10)

$$30 SV/SV_{rest} < L (11)$$

5

25

35

where L is a constant typically equal to a value of 1.2 to 1.5, depending on the health of the patient's myocardium.

Thus the lower pacing rate limit is continuously automatically calculated from the measured actual stroke volume SV and known values of SV_{rest} , HR_{rest} and the constant L. The actual stroke volume can be determined from e.g. bioimpedance measurements as will be described below.

Figure 1 is a block diagram of an embodiment of the pacemaker 10 according to the invention comprising a bioimpedance measurement unit 2 for measuring the time variation of the electric intracardiac bioimpedance $Z_{c}\left(t\right)$. This type of measurements is well known, see e.g. "Design of Cardiac Pacemakers", edited by John G. Webster, IEEE Press, 1995, pp. 380-386 and US-A-15 5,154,171, 5,280,429, 5,282,840 and 5,807,272. Thus the time variation of the intracardiac bioimpedance can be measured between the tip 4 of the intracardiac electrode 6 and the housing 8 of the pacemaker, when an excitation current is fed 20 from the electrode tip 4, as schematically illustrated in figure 2. Thus a standard pacing lead can be used for this measurement.

From the measured time variations $\Delta Z_c(t)$ the stroke volume SV needed for calculating the lower pacing rate limit according to equation (8) above, or for checking the inequalities (10) or (11), are determined in computing means 10, see figure 1.

The calculated lower limit value is supplied to a lower limit setting means 12 of a pacing rate limiter 14.

A pacing rate controller 16 is also provided for controlling the pacing rate of the pacer or pulse generator 18 in response to the patient's demands. In a limiting unit 20 of the limiter 14 the demanded pacing rate is compared to the set lower limit pacing rate and the actual pacing rate is limited to the set lower limit value if the demanded pacing WO 00/57953 PCT/SE00/00572

6

rate reaches this limit value. Thus in the pacemaker according to the invention a lower limit value for the pacing rate is continuously automatically determined and it is continuously automatically verified that the actual pacing rate does not exceed the present lower limit value.

Alternatively, the pacemaker can be modified to continuously monitor that the inequalities (10) or (11) above are satisfied.

10

15

5

Above bioimpedance measurements are described for determining the stroke volume SV. This parameter can, however, also be determined by other techniques, like by ECG measurements, by ultrasound technique, by radiometric and optical techniques etc. Generally all dynamic distance and/or capacity measuring methods are applicable.

PCT/SE00/00572

Claims

25

30

35

- 1. A rate adaptive pacemaker comprising a means for determining the demand of the patient's organism, a pacing rate controlling means for controlling the pacing rate in response to the patient's demand, and a pacing rate limiting means for preventing the pacing rate from becoming too low, characterized in that said pacing rate limiting means is adapted to limit the pacing rate downwards such that a first predetermined relation is satisfied between actual cardiac output (CO) and cardiac output (CO_{rest}) for the patient in rest conditions and a second predetermined relation is satisfied between actual stroke volume (SV_{rest}).
- 15 2. The pacemaker according to claim 1, characterized in that said first predetermined relation is

CO > CO_{rest}

and said second predetermined relation is

 $(SV)/(SV_{rest}) < L$

- where L denotes a predetermined constant > 1, preferably equal to a value between 1.2 and 1.5.
 - The pacemaker according to claims 1 or 2, characterized in that said pacing rate limiting includes a lower limit setting means for setting a lower limit value for the pacing rate, and a lower determining means for determining the relation between actual cardiac output (CO) and cardiac output (CO_{rest}) for the patient in rest conditions and the relation between actual stroke volume (SV) and a rest stroke volume (SV_{rest}) and calculating a lower pacing rate limit value from said relations for supply to said limit setting means.
 - 4. The pacemaker according to claim 3, characterized in that said lower limit determining means includes a stroke volume measuring means for measuring actual stroke volume SV and comparison means for comparing measured actual stroke volume SV with stroke volume SV $_{\rm rest}$ for the patient in rest conditions to ensure that the inequality

5

10

15

20

PCT/SE00/00572

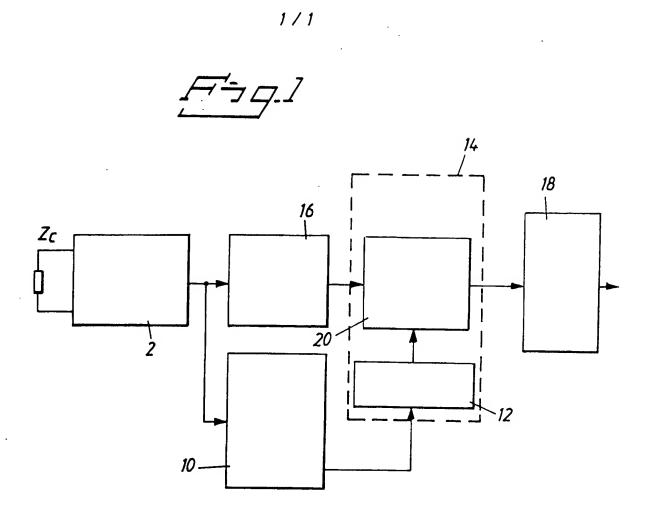
$SV/SV_{rest} < L$

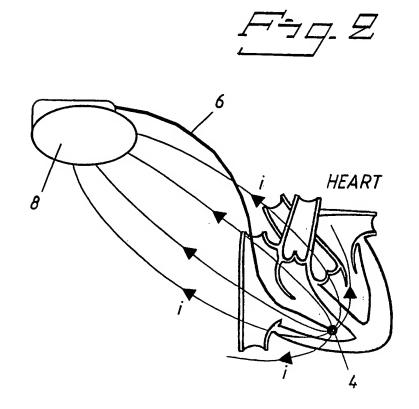
is satisfied, and in that said lower limit determining means is adapted to calculate a lower pacing rate limit value from the equation

lower pacing rate limit = HR_{rest} · (SV_{rest}/SV)

where HR_{rest} denotes the heart rate for the patient in rest conditions, provided that said inequality is satisfied.

- 5. The pacemaker according to any of the claims 2 4, characterized in that a bioimpedance measurement unit is provided to measure the cardiac bioimpedance as a function of time for determining therefrom actual cardiac output (CO) and actual stroke volume (SV) from the measured cardiac bioimpedance.
- 6. The pacemaker according to any of the claims 2 4, characterized in that an ECG measuring and analyzing unit is provided to measure ECG and determine therefrom actual cardiac output (CO) and actual stroke volume (SV).
- 7. The pacemaker according to any one of claims 1-4, characterized in that a dynamic distance measuring and analyzing unit is provided to determine therefrom actual cardiac output (CO) and actual stroke volume (SV).





International application No.

PCT/SE 00/00572

A. CLASSIFICATION OF SUBJECT MATTER IPC7: A61N 1/365 According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC7: A61N Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched SE, DK, FI, NO classes as above Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Category* Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. A EP 0140472 A1 (MEDTRONIC, INC.), 8 May 1985 1-6 (08.05.85), page 11, line 7 - page 13, line 30 A US 5183040 A (TIBOR A. NAPPHOLZ ET AL), 1-6 2 February 1993 (02.02.93), column 19. line 62 - column 20, line 3 EP 0576114 A2 (TELECTRONICS N.V.), A 1-6 29 December 1993 (29.12.93), column 29, line 23 - line 54 Further documents are listed in the continuation of Box C. See patent family annex. Special categories of cited documents "I" later document published after the international filing date or priority "A" document defining the general state of the art which is not considered date and not in conflict with the application but cited to understand the principle or theory underlying the invention to be of particular relevance "E" erlier document but published on or after the international filing date "X" document of particular relevance: the claimed invention carnot be document which may throw doubts on priority claim(s) or which is cited to establish the publication case of another citation or other considered novel or cannot be considered to involve an inversive step when the document is taken alone special reason (as specified) "Y" document of particular relevance: the claimed invention cames be document referring to an oral disclosure, use, exhibition or other considered to involve an inventive step when the document:s combined with one or more other such documents, such commandian document published prior to the international filing date but later than the priority date claimed being obvious to a person skilled in the art "&" document member of the same patent family Date of the actual completion of the international search Date of mailing of the international search report 2000 -07- 2 4 **26 June 2000** Name and mailing address of the ISA/ Authorized officer **Swedish Patent Office** Box 5055, S-102 42 STOCKHOLM Nikolaj Hautaviita/Els Facsimile No. +46 8 666 02 86 Telephone No. + 46 8 782 25 00

INTERNATIONAL SEARCH REPORT

Information on patent family members

02/12/99

International application No.

PCT/SE 00/00572

Patent document cited in search report			Publication date	Patent family member(s)			Publication date
EP	0140472	A1	08/05/85	CA JP JP JP US	1243361 1701308 3068708 60034462 4535774	C B A	18/10/88 14/10/92 29/10/91 22/02/85 20/08/85
US	5183040	A	02/02/93	NON	E		
EP	0576114	A2	29/12/93	DE US	576114 5197467		28/07/94 30/03/93

Form PCT/ISA/210 (patent family annex) (July 1992)

This Page is Inserted by IFW Indexing and Scanning Operations and is not part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

BLACK BORDERS

IMAGE CUT OFF AT TOP, BOTTOM OR SIDES

FADED TEXT OR DRAWING

BLURRED OR ILLEGIBLE TEXT OR DRAWING

SKEWED/SLANTED IMAGES

COLOR OR BLACK AND WHITE PHOTOGRAPHS

GRAY SCALE DOCUMENTS

LINES OR MARKS ON ORIGINAL DOCUMENT

REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY

IMAGES ARE BEST AVAILABLE COPY.

☐ OTHER:

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.